

REMARKS

Claims 1-43 are pending in the present application.

The rejections of: (a) Claims 1-3 under 35 U.S.C. §102(b) over Town et al, (b) Claims 1-5, 9-11, 19-21, 25-27, and 42 under 35 U.S.C. §102(b) over Dormann et al, and (c) Claims 1-11, 19-27, and 42 under 35 U.S.C. §102(e) over La Rosa et al, are obviated by amendment.

Applicants submit that none of Town et al, Dormann et al, or La Rosa et al disclose or suggest a polynucleotide falling within the scope of the claimed invention. Specifically, the sequences disclosed by Town et al have 63.7% homology on a nucleotide sequence level and 71.4% homology on an amino acid level. The sequences disclosed by Dormann et al have 63.2% homology on a nucleotide sequence level and 71.1% homology on an amino acid level. Further, the sequence disclosed by La Rosa et al has 90.1% homology on an amino acid level. Evidence for the same is provided by the Sequence Alignments **submitted herewith**. In view of the foregoing, the claimed invention is not anticipated by the cited references.

Withdrawal of these grounds of rejection is requested.

The rejections of: (a) Claims 1-11, 19-27, and 42 under 35 U.S.C. §112, first paragraph (enablement), and (b) Claims 1-11, 19-27, and 42 under 35 U.S.C. §112, first paragraph (written description), are believed to be obviated by amendment.

Indeed, it is the current trend in U.S. patent examination to narrow the permissible scope of homologs when DNA or protein sequences are claimed. This case falls right in line with this trend. Nonetheless, Applicants wish to direct the Examiner's attention to a recent decision by the U.S. PTO's Board of Patent Appeals and Interferences (*Ex parte Bandman*,

enclosed herewith) in which the Board held that claims to amino acid sequences that are at least 95% homologous to the disclosed sequence are adequately described and enabled when the specification describes the nucleotide and amino acid sequences.

As in *Ex parte Bandman*, the present specification provides the amino acid sequence (SEQ ID NO: 2) and the polynucleotide encoding the same (i.e., SEQ ID NO: 1). Moreover, the claims specify the activity required for all proteins encoded by the claimed polynucleotide that fall within the scope thereof. Clearly if the Board finds that under similar circumstances to the present specification an amino acid sequence having at least 95% homology is adequately described and enabled, the certainly so too is the homology of the present application.

Further, with respect to the sufficiency of the disclosure for describing the claimed sequence, the Examiner's attention is directed to Example 14 of the Synopsis of Application of Written Description Guidelines which analyzes a situation where a claim covers a protein that is at least 95% identical to a disclosed sequence and has a specific function. In these guidelines, the Patent Office has concluded that such a claim is adequately described within the meaning of 35 U.S.C. § 112, first paragraph

There is actual reduction to practice of the single disclosed species. The specification indicates that the genus of proteins that must be variants of SEQ ID NO: 3 does not have substantial variation since all of the variants must possess the specified catalytic activity and must have at least 95% identity to the reference sequence, SEQ ID NO: 3. The single species disclosed is representative of the genus because all members have at least 95% structural identity with the reference compound and because of the presence of an assay which applicant provided for identifying all of the at least 95% identical variants of SEQ ID NO: 3 which are capable of the specified catalytic activity. One of skill in the art would conclude that applicant was in possession of the necessary common attributes possessed by the members of the genus.

Conclusion: The disclosure meets the requirements of 35 USC §112 first paragraph as providing adequate written description for the claimed invention.

As the specification adequately describes the sequences that at least 95% homologous to SEQ ID NO: 2, a polynucleotide that is at least 95% homologous to SEQ ID NO: 1, and the specification describes how one can test for the recited activity to readily determine whether the variants are capable of the specified catalytic activity. Therefore, the claims as presented herein are deemed to be fully described and enabled.

Withdrawal of these grounds of rejection is requested.

The rejection of Claims 1-11, 19-27, and 42 under 35 U.S.C. §112, second paragraph, is obviated by amendment.

With respect to “stringent conditions” this language has been deleted in favor of the homology values recited in page 13, lines 19-28 used to define the “stringent conditions”. The term “gene” has been replaced with “polynucleotide”. Claims 11, 27, and 42 have been amended to ensure that all essential steps are recited.

Applicants request withdrawal of this ground of rejection.

The rejection of Claims 1-2, 11, and 27 under 35 U.S.C. §101 is obviated by amendment.

Claims 1 and 2 have been amended to define the polynucleotide as being “isolated”.

Withdrawal of this ground of rejection is requested.

The objection to the specification is obviated by the amendment to the description of Figure 4 and the submission of the enclosed substitute Sequence Listing. Applicants **submit**

herewith a substitute Sequence Listing and a corresponding computer-readable Sequence Listing. The sequence information recorded in the corresponding computer-readable Sequence Listing is identical to the paper copy of the substitute Sequence Listing. Support for all of the sequences listed in the substitute Sequence Listing is found in the present application. No new matter is believed to have been introduced by the submission of the substitute Sequence Listing and the corresponding computer-readable Sequence Listing. The specification has also been amended to add sequence identifiers where necessary. Support for this amendment is provided by the originally filed specification and Sequence Listing.

Finally, the objection to the drawings is obviated in part by amendment and traversed in part.

To address the criticism in paragraphs 5 and 6 of the Office Action, Applicants have amended the description of Figures 6-8, 10, 11, 13, and 16. Therefore, this objection is believed to be moot.

In paragraph 7 of the Office Action, the Examiner alleges that Figure 17 fails to comply with 37 CFR 1.84(g) "because it is framed". Applicants disagree with this allegation by the Examiner. Fig. 17 shows the results of genomic Southern hybridization described in Example 8. The solid line in Figure 17 is not a "frame" as the Examiner alleges, but rather is an illustration of the outer boundary of the membrane to which the content of the electrophoretic gel was transferred. Thus, the solid line in Figure 17 is not a "frame", but rather a part of the illustration. In view of the foregoing, Applicants submit that Figure 17 is in compliance with 37 CFR 1.84(g) and that this ground of rejection should be withdrawn.

Also, in paragraph 7, the Examiner alleges that the molecular size markers are missing for Figure 17. At the outset, it should be noted that there is no requirement in U.S.

patent practice for an electrophoretic gel to contain molecular size markers. This is especially true where the description in the specification clearly explains the detail of what is illustrated in the Figure. In this case, the description in Example 8 (see pages 44-45) sufficiently describes Figure 17 and what is shown therein. Further, the Examiner should be mindful of the fact that Figure 17 shows the results of a Southern hybridization assay where the probe is the kanamycin-tolerant (NPT) gene region of pBI121 labeled with Alphas Direct. Thus, following hybridization and detection, any molecular size markers present in the original electrophoretic gel would not be detected. In view of the foregoing, Applicants submit that Figure 17 is proper and complete.

In paragraph 8, the Examiner alleges that the molecular size markers are missing for Figure 18. This allegation is incorrect as it is noted that the molecular size markers are flanking lanes 1 and 6. Thus, this objection is without merit.

Applicants request withdrawal of these grounds of objection.

Applicants submit that the present application is in condition for allowance. Early notification to this effect is respectfully requested.

Respectfully submitted,

OBLON, SPIVAK, McCLELLAND,
MAIER & NEUSTADT, P.C.



Stephen G. Baxter, Ph.D.
Registration No. 32, 884

Customer Number
22850

Tel: (703) 413-3000
Fax: (703) 413-2220
(OSMMN 08/03)

Vincent K. Shier, Ph.D.
Registration No. 50,552

[GENETYX-MAC : Nucleotide Sequence Homology Data]

Date : 2007.07.27

1st Nucleotide Sequence

File Name : PsUGE1
Sequence Size : 1094

2nd Nucleotide Sequence

File Name : Doremann and Bennig DNA
Sequence Size : 1356

Unit Size to Compare = 4
Pick up Location = 5

[63.2% / 1023 bp] INT/OPT.Score : < 1604/ 1836 >

```
1' ATG
61" TTTGTTCTTCTGTTGGTGGTGGTATCTAGTTTTCAAAGAATCGATTTTGCCAAGTGGGT
4' GCGATCGGCGGGGCGGAGGCCGGGGGAGGCGCGGGGGCCAGCGGCCGGAGCGTGCTG
121" TCTTCTTGATAACCTTTCTTCTTTGAAATGGGTTCTTCTGTGGAGCAGAACATTCTT
64' GTGACGGGCGGCGCGGGGTTTCATCGGCACGCACACGGCGCTGCGCCTGCTGGAGCAGGGC
181" GTTACTGGTGGTGCTGGCTTTATCGGGACGCATACTGTTGTTCAACTTCTCAAAGATGGT
124' TACGGCGTCACCGTCGTCGACAACTTCCACAACTCCGTCCCCGAGGCGCTCGAACGCGTC
241" TTTAAGGTTTCGATCATCGATAATTTTGATAACTCTGTTATCGAAGCTGTTGATAGAGTT
184' CGCCTCATCGCCGGGCCCCGCGCTCTCCGCCCCGCTCGACTTCATCCGGGGGGATCTGAGG
301" AGGGAGCTTGTTGGTCCTGATCTCTCCAAGAAGCTCGACTTCAATCTGGGTGATCTAAGA
244' AGCGCCGGGGACTTGGAGAAGGCGTTTCGCGGCCAGGAGGTACGACGCGCTCGTCCACTTC
361" AACAAAGGGGACATTGAGAACTATTCTCCAAGCAGAGATTTGATGCTGTGATTCATTTT
304' GCGGGGCTCAAGGCCGTCGGGGAGAGCGTCGCGCGCCCGGACATGTACTACGAGAACAAC
421" GCGGGTCTTAAAGCTGTGGGTGAGAGTGTTGAAAAGGGTCGCCGCTACTTTGACAATAAC
364' CTCGCCGGCACCATCAACCTCTACAAGGCCATGAACGAGCACGGCTGCAAGAAGATGGTG
481" TTGGTTGGAACAATCAATCTATATGAGACCATGGCAAAGTACAACTGCAAAATGATGGTG
424' TTCTCGTCGTCCGCGACCGTGACGGCTGGCCGGAGGTGATCCCGTGCGTCGAGGACTCC
541" TTTTCATCTTCTGCCACTGTTTATGGACAACCTGAAAAGATTCCATGCATGGAAGACTTT
484' AAGCTGCAGGCCGCCAACCCTACGGCAGGACCAAGCTCATCCTGGAGGAGTTGGCGCGG
601" GAATTAAAGGCTATGAATCCTTATGGTCGTACTAAGCTCTTTCTTGAAGAAATAGCTAGA
544' GACTACCAGCGCGCGGACCCGGGCTGGAGCATCGTCCTGCTGCGCTACTTCAACCCCATC
661" GATATTCAAAGGCAGAACCGGAATGGAGAATTATTCTGCTGAGGTAATTCAATCCTGTA
604' GGCGCCACAGCTCCGGCGAGATCGGCGAGGACCCCAAGGGGGTGCCCAACAACCTGCTG
721" GGAGCACATGAGAGTGGCAGTATTGGTGAGGATCCAAAAGGCATCCCCAATAACCTCATG
664' CCCTACATCCAGCAGGTCGCCGTCGGCAGGCTCCCCGAGCTCAACGTCTACGGCCACGAT
781" CCTTACATCCAACAAGTGGCCGTTGGACGTTTACCGGAACCTCAATGTCTATGGACATGAC
```


306' GGGGCTCAAGGCCGTCGGGGAGAGCGTCGCGCGCCCGGACATGTACTACGAGAACAACCT
 *** **
 481" GGGTCTTAAAGCTGTGGGTGAGAGTGTTGAAAACCTCGCCGCTACTTTGACAATAACTT
 366' CGCCGGCACCATCAACCTCTACAAGGCCATGAACGAGCACGGCTGCAAGAAGATGGTGTT
 * **
 541" GGTGGAACAATCAATCTATATGAGACCATGGCAAAGTACAACGCAAAATGATGGTGTT
 426' CTCGTCGTCCGCGACCGTGACGGCTGGCCGGAGGTGATCCCGTGCGTCGAGGACTCCAA
 *
 601" TTCATCTTCTGCCACTGTTTATGGACAACCTGAAAAGATTCCATGCATGGAAGACTTTGA
 486' GCTGCAGGCCGCAACCCCTACGGCAGGACCAAGCTCATECTGGAGGAGTTGGCGCGGGA
 *
 661" ATTAAGGCTATGAATCCTTATGGTCGTAAGCTCTTTCTTGAAGAAATAGCTAGAGA
 546' CTACCAGCGCGCGGACCCGGGCTGGAGCATCGTCCTGCTGCGCTACTTCAACCCCATCGG
 *
 721" TATTCAAAGGCAGAACCGGAATGGAGAATTATTCTGCTGAGGTAATTCAATCCTGTAGG
 606' CGCCACAGCTCCGGCGAGATCGGCGAGGACCCCAAGGGGGTGCCCAACAACCTGCTGCC
 *
 781" AGCACATGAGAGTGGCAGTATTGGTGAGGATCCAAAAGGCATCCCCAATAACCTCATGCC
 666' CTACATCCAGCAGGTGCGCGTCGGCAGGCTCCCCGAGCTCAACGTCTACGGCCACGATTA
 *
 841" TTACATCCAACAAGTGGCCGTTGGACGTTTACCGGAACCTCAATGTCTATGGACATGACTA
 726' CCCCACCCGTGACGGCACCGCGATCAGGGACTACATACACGTCGTCGACCTGGCCGACGG
 *
 901" TCCCACCGAGGATGGTAGTGCGGTAAGAGACTACATCCATGTGATGGATTTAGCAGATGG
 786' GCACATCGCGGCGCTGAACAAGCTGTTGACACTCCTGATTTGCGTTGTGTGGCCTACAA
 *
 961" CCATATCGCTGCGCTCAGGAAGCTATTTGCTGATCCAAAGATTGGTTGTACTGCTTACAA
 846' TCTGGGCACAGGGCGCGGCACATCCGTTCTCGAGATGGTGGCGGCGTTCAAGAAGGCATC
 *
 1021" TCTAGGACTGGTCAAGGAACGTCTGTGTTAGAAATGGTTGCAGCTTTTGAAAAAGCTTC
 906' CGGCAAGGAGATCCCCACCAAGATGTGCCCCAGGAGACCGGGTGACGCGACGGAGGTTTA
 *
 1081" CGGCAAGAAAATCCCGATTAAGCTCTGTCCGAGAAGGTCAGGAGATGCAACAGCAGTTTA
 966' CGCGTCCACTGAGAAGGCCGAAAGGGAGCTCGGATGGAGGGCCCAGTATGGAATCGAGGA
 *
 1141" TGCTTCAACAGAGAAGGCTGAGAAAGAACTTGGCTGGAAGGCAAAATATGGAGTGGATGA
 1026' GATGTGCAGGGACCAAGTGGAACTGGGCCAAGAAGAACCCTATGGCTACTGCGGCACTGC
 *
 1201" GATGTGCAGAGATCAGTGGAAATGGGCAAACAATAATCCATGGGGTTACCAGAATAAGCT
 1086' CGAAAAATA
 1261" TTGAATTTACTTCTTTTGTGGAGTTACCATTTCTAATTACTCAAATCTAAAAGAAAGA

[GENETYX-MAC: Multiple Alignment]

Date : 2007.07.26

Doremann and Bennig DNA	1	-----GA	2
Town et al UGE DNA	1	1 CCACATCATTTTCTATTTTTCGCTTTCGCTTCTTATCAACTTGTAAACAAAGCTA	60
PsUGE1	1	-----	1
Doremann and Bennig DNA	3	CAAAATATCTTTAAATAAGGACCCAACCTCTTTTCAATTCCTCCCATCAATCTTCTTATT	62
Town et al UGE DNA	61	CAAAATATCTTTAAATAAGGACCCAACCTCTTTTCAATTCCTCCCATCAATCTTCTTATT	120
PsUGE1	1	-----	1
Doremann and Bennig DNA	63	TGTTCTTCTGTTGGTGGTGGTATCTAGTTTTCAAAGAATCGATTTTGCCAAGTGGGTTG	122
Town et al UGE DNA	121	TGTTCTTCTGTTGGTGGTGGTATCTAGTTTTCAAAGAATCGATTTTGCCAAGTGGGTTG	180
PsUGE1	1	-----ATGGCGA---	7
Doremann and Bennig DNA	123	TTCTTGATAACCTTTCTTCTTTTGAATGGGTTCTT-CTGTGGAGCAGAACATTCTTG	181
Town et al UGE DNA	181	TTCTTGATAACCTTTCTTCTTTTGAATGGGTTCTT-CTGTGGAGCAGAACATTCTTG	239
PsUGE1	8	TCGGCGGGGCGGAGG--GGGGGGGGGGCGGGGGCGAGCGCGG-GAGGCTGCTGG	64
Doremann and Bennig DNA	182	TTACTGGTGGTGGTGGCTTTATCGGGACGCATACTGTTGTTCAACTTCTCAAAGATGGTT	241
Town et al UGE DNA	240	TTACTGGTGGTGGTGGCTTTATCGGGACGCATACTGTTGTTCAACTTCTCAAAGATGGTT	299
PsUGE1	65	TGACGGGCGGCGGGGCTTCTATCGGCACGCACAGCGCGTGGCGCTGCTGGAGCGGGCT	124
Doremann and Bennig DNA	242	TTAAGGTTTCGATCATCGATAATTTTGATAACTCTGTTATCGAAGCTGTTGATAGAGTTA	301
Town et al UGE DNA	300	TTAAGGTTTCGATCATCGATAATTTTGATAACTCTGTTATCGAAGCTGTTGATAGAGTTA	359
PsUGE1	125	ACGGGCTCAGCGTGGTGGGAACTTCCCAACTCGTCCCGAGGCGGCTCAACGCTCC	184
Doremann and Bennig DNA	302	GGGAGCTTGTGGTCTGATCTCTCAAGAAGCTCGACTTCAATCTGGGTGATCTAAGAA	361
Town et al UGE DNA	360	GGGAGCTTGTGGTCTGATCTCTCAAGAAGCTCGACTTCAATCTGGGTGATCTAAGAA	419
PsUGE1	185	GCTCAATCCCGCGGCGGCTCTCGGCGCCCGCTCGACTTCACTGGGGGATCTGAGGA	244
Doremann and Bennig DNA	362	ACAAAGGGGACATTGAGAACTATTCTCAAGCAGAGATTGATGCTGTGATTCAATTTTG	421
Town et al UGE DNA	420	ACAAAGGGGACATTGAGAACTATTCTCAAGCAGAGATTGATGCTGTGATTCAATTTTG	479
PsUGE1	245	GGCCGGGGGATTTGGAGAAAGCGTTGGGGCCAGGAGGACGACGCGCTCGTTCACCTCG	304
Doremann and Bennig DNA	422	CGGGTCTTAAAGCTGTGGGTGAGAGTGTGAAAAGGGTCGCCGCTACTTTGACAATACT	481
Town et al UGE DNA	480	CGGGTCTTAAAGCTGTGGGTGAGAGTGTGAAAAGGGTCGCCGCTACTTTGACAATACT	539
PsUGE1	305	CGGGCTCAAGGCGCTGGGAGAGCTCGCGCGCGGAGATGTAATACGAGAAACAC	364
Doremann and Bennig DNA	482	TGGTTGGAACAATCAATCTATATGAGACCATGGCAAAGTACAACCTGCAAATGATGGTGT	541
Town et al UGE DNA	540	TGGTTGGAACAATCAATCTATATGAGACCATGGCAAAGTACAACCTGCAAATGATGGTGT	599
PsUGE1	365	TCGCGCGCAGCTCAACTCTTACACAGGCTCATGAACGAGCAGGGCTGCAAGGAGATGGTGT	424
Doremann and Bennig DNA	542	TTTCATCTTCTGCCACTGTTTATGGACAACCTGAAAAGATTCCATGCTGGAAGACTTTG	601
Town et al UGE DNA	600	TTTCATCTTCTGCCACTGTTTATGGACAACCTGAAAAGATTCCATGCTGGAAGACTTTG	659
PsUGE1	425	TCTGTGTGTGCGGAGCTGTATCGGCTGGCGGAGGTGATCTGCTGGTTCAGGACTTCCA	484
Doremann and Bennig DNA	602	AATTAAAGGCTATGAATCCTTATGGTCGTAAGCTCTTTCTTGAAGAAATAGCTAGAG	661
Town et al UGE DNA	660	AATTAAAGGCTATGAATCCTTATGGTCGTAAGCTCTTTCTTGAAGAAATAGCTAGAG	719
PsUGE1	485	AGCTTGCAGGCGCCAACTCTACGCGAGGACCAAGCTCACTTGGAGGATTTGGCGCGG	544
Doremann and Bennig DNA	662	ATATTCAAAAGGCAGAACCGGAATGGAGAATTATCTGCTGAGGTACTTCAATCTGTAG	721
Town et al UGE DNA	720	ATATTCAAAAGGCAGAACCGGAATGGAGAATTATCTGCTGAGGTACTTCAATCTGTAG	779
PsUGE1	545	ACTACAGCGCGCGGACCGGCGTGGAGCATCTGCTGCTGCTACTTCAACCTCACTCG	604
Doremann and Bennig DNA	722	GAGCACATGAGAGTGGCAGTATTGGTGAGGATCCAAAAGGCATCCCAATAACCTCATGC	781
Town et al UGE DNA	780	GAGCACATGAGAGTGGCAGTATTGGTGAGGATCCAAAAGGCATCCCAATAACCTCATGC	839
PsUGE1	605	CGGCTACAGCTCCGGGAGATCGGCGAGGACCGCAAGGGGGTGGCCAAACACCTGCTGC	664
Doremann and Bennig DNA	782	CTTACATCCAACAAGTGGCGTGGACGTTTACCGGAACCTCAATGCTATGGACATGACT	841
Town et al UGE DNA	840	CTTACATCCAACAAGTGGCGTGGACGTTTACCGGAACCTCAATGCTATGGACATGACT	899
PsUGE1	665	CTTACATCCAACAAGTGGCGTGGACGTTTACCGGAACCTCAATGCTATGGACATGACT	724
Doremann and Bennig DNA	842	ATCCCACCGAGGATGGTAGTGCGGTAAAGAGACTACATCCATGTGATGGATTTAGCAGATG	901
Town et al UGE DNA	900	ATCCCACCGAGGATGGTAGTGCGGTAAAGAGACTACATCCATGTGATGGATTTAGCAGATG	959
PsUGE1	725	ACCCACCGTGAAGGACCGCGGATGAGGACTACATACAGCTGCTGACCTGGGCGAGCG	784
Doremann and Bennig DNA	902	GCCATATCGCTGCGCTCAGGAAGCTATTTGCTGATCCAAAGATTGGTTGACTGCTTACA	961
Town et al UGE DNA	960	GCCATATCGCTGCGCTCAGGAAGCTATTTGCTGATCCAAAGATTGGTTGACTGCTTACA	1019
PsUGE1	785	GCACATCGGCGCGCTGAGCAAGCTGTTCACACTCTGTATTTGGTTGTGTGCTTACA	844
Doremann and Bennig DNA	962	ATCTAGGGACTGGTCAAGGAACGTCTGTGTTAGAAATGGTTGCAGCTTTTGAAAAGCTT	1021
Town et al UGE DNA	1020	ATCTAGGGACTGGTCAAGGAACGTCTGTGTTAGAAATGGTTGCAGCTTTTGAAAAGCTT	1079
PsUGE1	845	ATCTAGGGACTGGTCAAGGAACGTCTGTGTTAGAAATGGTTGCAGCTTTTGAAAAGCTT	904
Doremann and Bennig DNA	1022	CCGGCAAGAAAATCCCGATTAAAGCTCTGTCCGAGAAGGTCAGGAGATGCAACAGCAGTTT	1081
Town et al UGE DNA	1080	CCGGCAAGAAAATCCCGATTAAAGCTCTGTCCGAGAAGGTCAGGAGATGCAACAGCAGTTT	1139
PsUGE1	905	CCGGCAAGGATCCCGCTCCCAAGATGTGCTCAGGAGACGGGTGACGGGACGGAGGTTT	964
Doremann and Bennig DNA	1082	ATGCTTCAACAGAGAAGGCTGAGAAAGAACTTGGCTGGAAGGCAAAATATGGAGTGGATG	1141
Town et al UGE DNA	1140	ATGCTTCAACAGAGAAGGCTGAGAAAGAACTTGGCTGGAAGGCAAAATATGGAGTGGATG	1199
PsUGE1	965	ACGGGTCACTGAGAAAGGCGAAAGGGAGCTCGATGGAGGGCCAGTATGGAATCGAGG	1024
Doremann and Bennig DNA	1142	AGATGTGCAGAGATCAGTGGAAATGGGCTTTCAATAATCCATGGGGTTACAGAATAAGC	1201
Town et al UGE DNA	1200	AGATGTGCAGAGATCAGTGGAAATGGGCTTTCAATAATCCATGGGGTTACAGAATAAGC	1259
PsUGE1	1025	AGATGTGCAGAGATCAGTGGAAATGGGCTTTCAATAATCCATGGGGTTACAGAATAAGC	1084
Doremann and Bennig DNA	1202	TTTGAATTTACTTCTTTTGGTGGAGTTACCATTTCTAATTACTCAAATCTAAAAGAAAG	1261
Town et al UGE DNA	1260	TTTGAATTTACTTCTTTTGGTGGAGTTACCATTTCTAATTACTCAAATCTAAAAGAAAG	1319
PsUGE1	1085	CCGAAATAA-----	1094
Doremann and Bennig DNA	1262	AAATATACATACATATGATGATATAGTTGTGCTTTATATTCACATGTATCGAACTGATG	1321
Town et al UGE DNA	1320	AAATATACATACATATGATGATATAGTTGTGCTTTATATTCACATGTATCGAACTGATG	1379
PsUGE1	1094	-----	1094
Doremann and Bennig DNA	1322	TCTTACTTCGATGAATAAAATGGAAGTTGATTTA-----	1356
Town et al UGE DNA	1380	TCTTACTTCGATGAATAAAATGGAAGTTGATTTGATTGAATTTTATGTTTCTTTCACT	1439
PsUGE1	1094	-----	1094
Doremann and Bennig DNA	1356	-----	1356
Town et al UGE DNA	1440	GAATAAAAGGCTTGTTCATGG	1462
PsUGE1	1094	-----	1094

[GENETYX-MAC : Amino Acid Sequence Homology Data]

Date : 2007.07.27

1st Amino Acid Sequence

File Name : PsUGE1TRANSLATE
Sequence Size : 364

2nd Amino Acid Sequence

File Name : Doremann bennig translate
Sequence Size : 351

Unit Size to compare = 2
Pick up Location = 5

[71.1% / 346 aa]

INT/OPT.Score : < 1348/ 1356 >

```
1' MAIGGAEAGGGGAGASGRSVLVTGGAGFIGHTHTALRLLEQGYGVTVVDNFHNVSYPEALER
*****
1" MGSSVEQNILVTGGAGFIGHTHTVVQLKDGFKVSIIDNFDNSVIEAVDR

61' VRLIAGPALSARLDFIRGDLRSAGDLEKAFAARRYDAVVHFAGLKAVGESVARPDMYYEN
** ** ** ** **
50" VRELVGPDLSKKLDFNLGDLRNKGDIEKLFSKQRFDAVIHFAGLKAVGESVEKGRRYFDN

121' NLAGTINLYKAMNEHGCKKMFSSSATVYGWPEVIPCVEDSKLQAANPYGRTKLILEELA
** ** ** ** **
110" NLVGTINLYETMAKYNCKMMVFSSSATVYGQPEKIPCMEDFELKAMNPYGRTKLFLEEIA

181' RDYQRADPGWSIVLLRYFNPIGAHSSGEIGEDPKGVPNNLLPYIQQVAVGRLPELNVYGH
** ** ** ** **
170" RDIQKAPEWRIILLRYFNPVGAHESGSIGEDPKGIPNNLMPYIQQVAVGRLPELNVYGH

241' DYPTRDGTAIRDYIHVVDLADGHIAALNKLFDTPDFGCVAYNLGTGRGTSVLEMVAAFKK
**** ** ** ** **
230" DYPTEDGSAVRDYIHVMDLADGHIAALRKLFAADPKIGCTAYNLGTGQTSVLEMVAAFEK

301' ASGKEIPTKMCPRRPGDATEVYASTEKAERELGWRAQYGIEEMCRDQWNWAKKNPYGYCG
**** ** ** ** **
290" ASGKKIPIKLCPRRSGDATAVYASTEKAEEKELGWKAKYGVDEMCRDQWKWAFNNPWGYQN

361' TAEK

350" KL
```

1st Amino Acid Sequence

File Name : PsUGE1TRANSLATE
Sequence Size : 364

2nd Amino Acid Sequence

File Name : Town et al. UGE
Sequence Size : 351

Unit Size to compare = 2
Pick up Location = 5

[71.4% / 346 aa]

INT/OPT.Score : < 1358/ 1366 >

```
1' MAIGGAEAGGGGAGASGRSVLVTGGAGFIGHTHTALRLLEQGYGVTVVDNFHNVSYPEALER
*****
1" MGSSVEQNILVTGGAGFIGHTHTVVQLKDGFKVSIIDNFDNSVIEAVDR
```


[GENETYX-MAC: Multiple Alignment]

Date : 2007.07.27

PsUGE1amino acids	1	MAIGGAEAGGGGAGASGRSVLVTGGAGFIGHTHTALRLLEQGYGNTVVVDFHNSVPEALER	60
Town et al. UGE amino acids	1	-----MGS---SVEQN-ILVTGGAGFIGHTHTVQLLKDGFKMSIIDNFNSVIEAVDR	49
Doremann& Bennig amino acids	1	-----MGS---SVEQN-ILVTGGAGFIGHTHTVQLLKDGFKMSIIDNFNSVIEAVDR	49
Rosa Patent amino acids	1	MAIGGSEAGGGGAGSMR-SVLVTGGAGFIGHTHTVLRLLLEQGTIMTVVDFHNSVPEALDR	59
PsUGE1amino acids	61	VRL-IAGPALSARLDIFIRGDLRSAGDLEKAF AARRYDAVVF-FAGLKAVGESVARPDMYY	118
Town et al. UGE amino acids	50	VRELV-GPDL SKKLD FNLGDLRNKGDI EKLF SKQRFDAVIH-FAGLKAVGESVENPRRMF	107
Doremann& Bennig amino acids	50	VRELV-GPDL SKKLD FNLGDLRNKGDI EKLF SKQRFDAVIH-FAGLKAVGESVEKGRRMF	107
Rosa Patent amino acids	60	VRL-IAGPALSTRLDIFIRGDLRNTDLEKVF AARRYDAVIHPFAGLKAVGESVAHPDMYY	118
PsUGE1amino acids	119	ENNLAGTINLYKAMNEHGCKKMFSSSATVYQWPEVIPCVEDSKLQAANPYGRTKLILEE	178
Town et al. UGE amino acids	108	DNNLVGTINLYETMAKYNCKMMVFSSSATVYGOPEKIPCMEDFELKAMNPYGRTKLFL EE	167
Doremann& Bennig amino acids	108	DNNLVGTINLYETMAKYNCKMMVFSSSATVYGOPEKIPCMEDFELKAMNPYGRTKLFL EE	167
Rosa Patent amino acids	119	ENNLI GTINLYKSMKEHGCKKLVFSSSATVYQWPEVIPCVEDSKLQAANPYGRTKLILED	178
PsUGE1amino acids	179	LARDYQRADPGNSIVLLRYFNPIGAHSSGEIGEDPKGIPNNLLPYIQQVAVGRLPELNVY	238
Town et al. UGE amino acids	168	IARDIQKAEPWRDILLRYFNPIGAHSSGEIGEDPKGIPNNLLPYIQQVAVGRLPELNVY	227
Doremann& Bennig amino acids	168	IARDIQKAEPWRDILLRYFNPIGAHSSGEIGEDPKGIPNNLLPYIQQVAVGRLPELNVY	227
Rosa Patent amino acids	179	MARDYHRADTEWSIVLLRYFNPIGAHSSGEIGEDPKGIPNNLLPYIQQVAVGRAPXNLNVY	238
PsUGE1amino acids	239	GHDYPTRDGTAIRDYIHVMDLADGHIAALNKLFDTPDFGQVAYNLGTGRGTSVLEMVAAF	298
Town et al. UGE amino acids	228	GHDYPTEDGSAVRDYIHVMDLADGHIAALRKLHADPKIGCTAYNLGTGQGTSVLEMVAAF	287
Doremann& Bennig amino acids	228	GHDYPTEDGSAVRDYIHVMDLADGHIAALRKLHADPKIGCTAYNLGTGQGTSVLEMVAAF	287
Rosa Patent amino acids	239	GHDYPTRDGTAIRDYIHVMDLADGHIAALKKLFDSPDITGVAYNLGTGRGTSVLEMVAAF	298
PsUGE1amino acids	299	KKASGKEIPTKMCPRRPGDATEVYASTEKAERELGWRAQYGIEMCRDQWNWAKKNPYG-	357
Town et al. UGE amino acids	288	EKASGKKIPIKLCPRRSGDATAVYASTEKAERELGWKAKYGVDEMCRDQWKWANNNPWGY	347
Doremann& Bennig amino acids	288	EKASGKKIPIKLCPRRSGDATAVYASTEKAERELGWKAKYGVDEMCRDQWKWAFNNPWGY	347
Rosa Patent amino acids	299	KKASGKEIPTKLCPRRPGDAT-EVYASTEKAERELAWRAQYGIEMCRDQWNWAKKNPYG-	356
PsUGE1amino acids	358	--YCGTAEK	364
Town et al. UGE amino acids	348	QNKL-----	351
Doremann& Bennig amino acids	348	QNKL-----	351
Rosa Patent amino acids	357	--YCGGAKK	363